

10/534,945

=> file caplus

FILE 'CAPLUS' ENTERED AT 16:34:21 ON 28 MAY 2008

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FILE COVERS 1907 - 28 May 2008 VOL 148 ISS 22

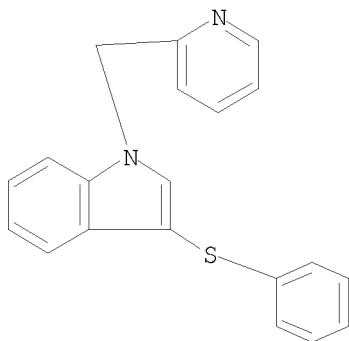
FILE LAST UPDATED: 27 May 2008 (20080527/ED)

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<http://www.cas.org/legal/infopolicy.html>

=> d que

L1 STR



Structure attributes must be viewed using STN Express query preparation.

L3 30 SEA FILE=REGISTRY SSS FUL L1

L4 2 SEA FILE=CAPLUS L3

=> d l4 1-2 ibib abs hitstr

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2005:1170665 CAPLUS

DOCUMENT NUMBER: 143:440257

TITLE: Preparation of indole derivatives as androgens

INVENTOR(S): Van Der Louw, Jaap; Teerhuis, Neeltje Miranda;
Lommerse, Johannes Petrus Maria; Stock, Herman Thijs;
Hermkens, Pedro Harold Han

PATENT ASSIGNEE(S): Akzo Nobel N.V., Neth.

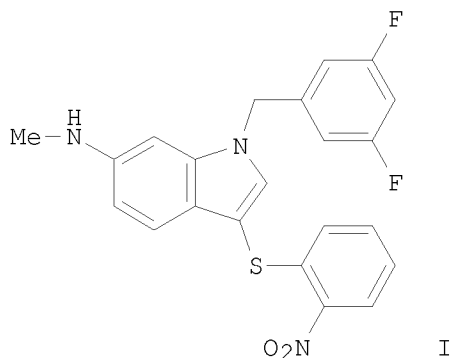
SOURCE: PCT Int. Appl., 65 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005102998	A1	20051103	WO 2005-EP51766	20050421
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
EP 1589003	A1	20051026	EP 2004-101700	20040423
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK				
AU 2005235751	A1	20051103	AU 2005-235751	20050421
CA 2562571	A1	20051103	CA 2005-2562571	20050421
EP 1742912	A1	20070117	EP 2005-737941	20050421
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, HR				
CN 1956954	A	20070502	CN 2005-80016406	20050421
BR 2005010078	A	20071016	BR 2005-10078	20050421
JP 2007533707	T	20071122	JP 2007-508908	20050421
MX 2006PA12201	A	20070117	MX 2006-PA12201	20061020
US 20070225352	A1	20070927	US 2006-587192	20061020
PRIORITY APPLN. INFO.:			EP 2004-101700	A 20040423
			US 2004-565043P	P 20040423
			WO 2005-EP51766	W 20050421
OTHER SOURCE(S):		CASREACT 143:440257; MARPAT 143:440257		
GI				



AB Indole derivs. were prepared and tested for androgenic activity. E.g., I was prepared starting from 6-nitroindole and 1-bromomethyl-3,5-difluorobenzene. The preparation of a number of indole derivs. was given and extensive androgenic activity data.

IT 868672-40-6P 868672-63-3P 868672-70-2P

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868672-71-3P 868672-74-6P 868672-78-0P

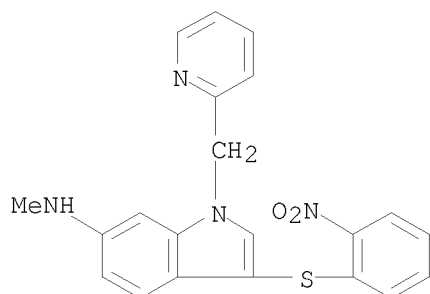
868672-79-1P 868672-85-9P 868672-86-0P

RL: PAC (Pharmacological activity); PRP (Properties); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of indole derivs. as androgens)

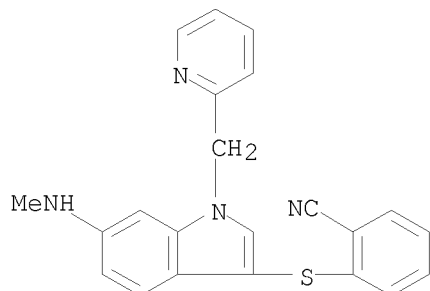
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CN 1H-Indol-6-amine, N-methyl-3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)-
(CA INDEX NAME)



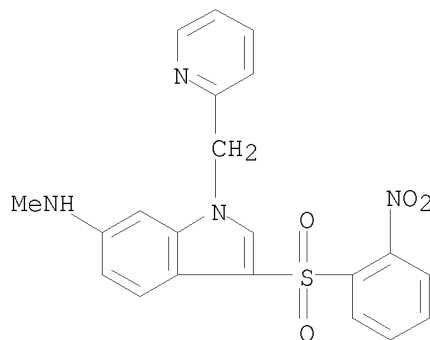
RN 868672-63-3 CAPLUS

CN Benzonitrile, 2-[[6-(methyamino)-1-(2-pyridinylmethyl)-1H-indol-3-yl]thio]- (CA INDEX NAME)



RN 868672-70-2 CAPLUS

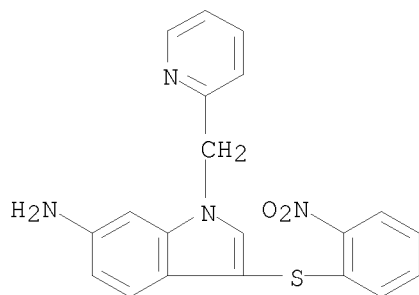
CN 1H-Indol-6-amine, N-methyl-3-[(2-nitrophenyl)sulfonyl]-1-(2-pyridinylmethyl)- (CA INDEX NAME)



10/534,945

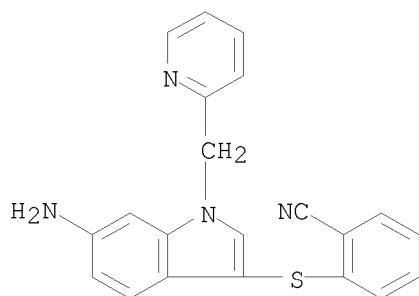
RN 868672-71-3 CAPLUS

CN 1H-Indol-6-amine, 3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)- (CA INDEX NAME)



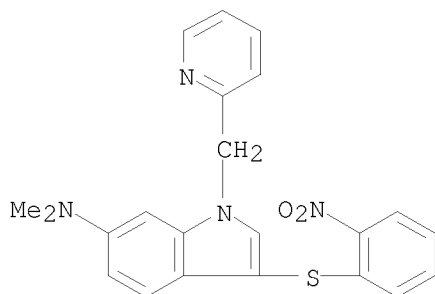
RN 868672-74-6 CAPLUS

CN Benzonitrile, 2-[[6-amino-1-(2-pyridinylmethyl)-1H-indol-3-yl]thio]- (CA INDEX NAME)



RN 868672-78-0 CAPLUS

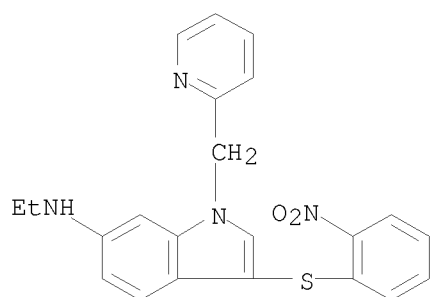
CN 1H-Indol-6-amine, N,N-dimethyl-3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)- (CA INDEX NAME)



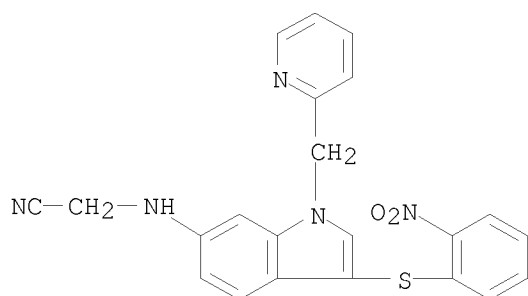
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CN 1H-Indol-6-amine, N-ethyl-3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)- (CA INDEX NAME)

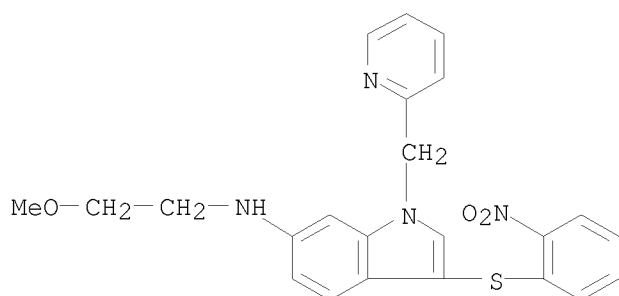
10/534,945



RN 868672-85-9 CAPLUS
CN Acetonitrile, 2-[[3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)-1H-indol-6-yl]amino]- (CA INDEX NAME)

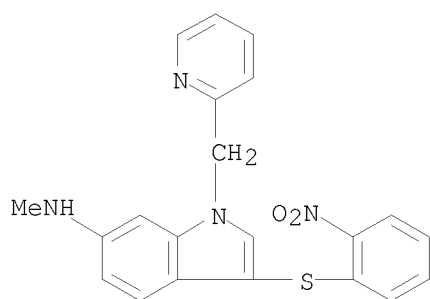


RN 868672-86-0 CAPLUS
CN 1H-Indol-6-amine, N-(2-methoxyethyl)-3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)- (CA INDEX NAME)



IT 868672-98-4P 868673-07-8P 868673-08-9P
868673-34-1P 868673-35-2P 868673-36-3P
868673-37-4P 868673-39-6P 868673-56-7P
RL: PRP (Properties); RCT (Reactant); SPN (Synthetic preparation); PREP
(Preparation); RACT (Reactant or reagent)
(preparation of indole derivs. as androgens)
RN 868672-98-4 CAPLUS
CN 1H-Indol-6-amine, N-methyl-3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)-,
hydrochloride (1:?) (CA INDEX NAME)

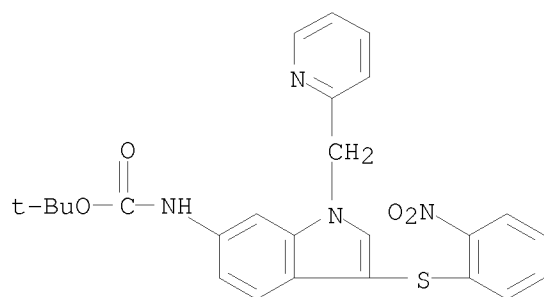
10/534,945



● x HCl

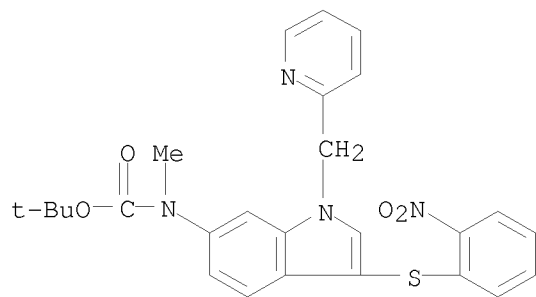
RN 868673-07-8 CAPLUS

CN Carbamic acid, [3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)-1H-indol-6-yl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



RN 868673-08-9 CAPLUS

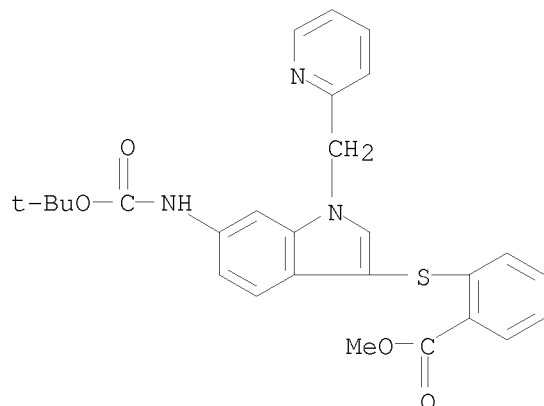
CN Carbamic acid, methyl[3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)-1H-indol-6-yl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



RN 868673-34-1 CAPLUS

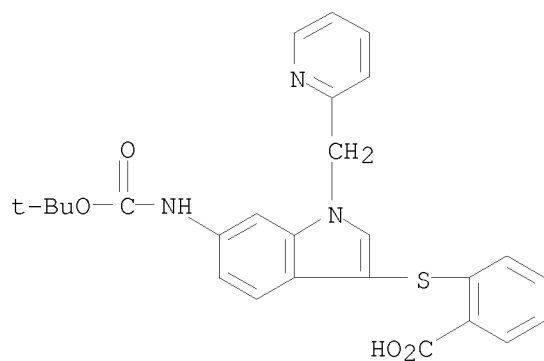
CN Benzoic acid, 2-[[6-[[[(1,1-dimethylethoxy)carbonyl]amino]-1-(2-pyridinylmethyl)-1H-indol-3-yl]thio]-, methyl ester (CA INDEX NAME)

10/534,945



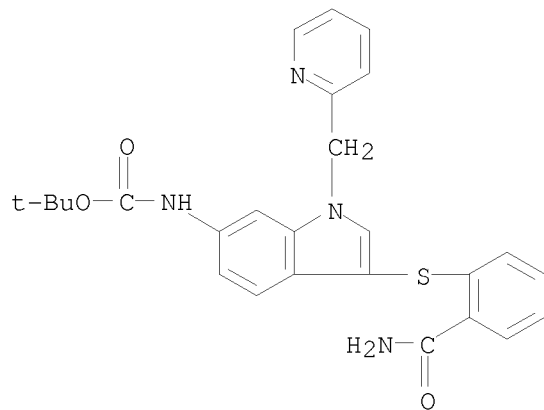
RN 868673-35-2 CAPLUS

CN Benzoic acid, 2-[[6-[[[(1,1-dimethylethoxy)carbonyl]amino]-1-(2-pyridinylmethyl)-1H-indol-3-yl]thio]- (CA INDEX NAME)



RN 868673-36-3 CAPLUS

CN Carbamic acid, [3-[[2-(aminocarbonyl)phenyl]thio]-1-(2-pyridinylmethyl)-1H-indol-6-yl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

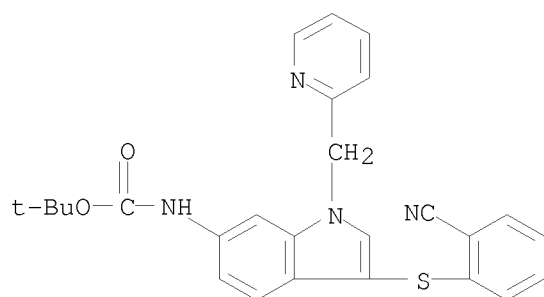


RN 868673-37-4 CAPLUS

CN Carbamic acid, [3-[(2-cyanophenyl)thio]-1-(2-pyridinylmethyl)-1H-indol-6-

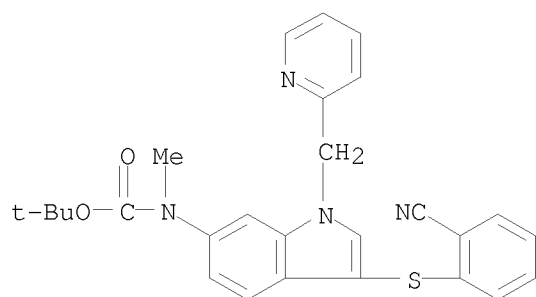
10/534,945

yl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



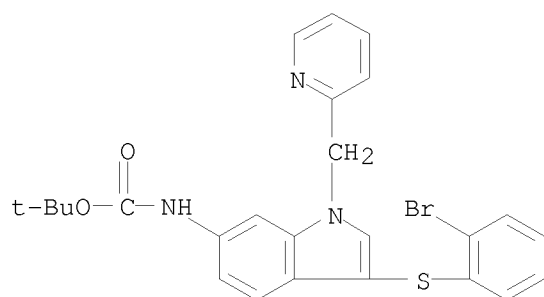
RN 868673-39-6 CAPLUS

CN Carbamic acid, [3-[(2-cyanophenyl)thio]-1-(2-pyridinylmethyl)-1H-indol-6-yl]methyl-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



RN 868673-56-7 CAPLUS

CN Carbamic acid, [3-[(2-bromophenyl)thio]-1-(2-pyridinylmethyl)-1H-indol-6-yl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2004:412918 CAPLUS

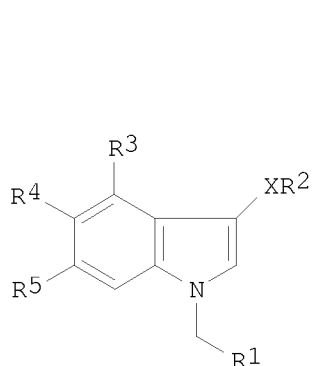
DOCUMENT NUMBER: 140:423584

TITLE: A preparation of indole derivatives useful in the treatment of androgen-receptor related diseases

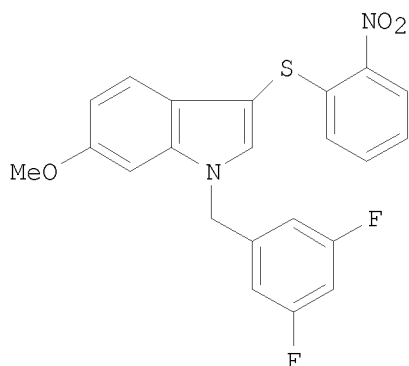
INVENTOR(S): Hermkens, Pedro Harold Han; Stock, Herman Thijs;
 Teerhuis, Neeltje Miranda; Lommerse, Johannes Petrus
 Maria; Van der Louw, Jaap
 PATENT ASSIGNEE(S): Akzo Nobel N.V., Neth.
 SOURCE: PCT Int. Appl., 75 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004041782	A1	20040521	WO 2003-EP50783	20031103
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2504907	A1	20040521	CA 2003-2504907	20031103
AU 2003301853	A1	20040607	AU 2003-301853	20031103
BR 2003016020	A	20050920	BR 2003-16020	20031103
EP 1585727	A1	20051019	EP 2003-810458	20031103
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
CN 1714078	A	20051228	CN 2003-80103950	20031103
JP 2006507293	T	20060302	JP 2004-549180	20031103
NO 2005002012	A	20050526	NO 2005-2012	20050425
ZA 2005003559	A	20060830	ZA 2005-3559	20050504
IN 2005CN00826	A	20070817	IN 2005-CN826	20050504
MX 2005PA04929	A	20050818	MX 2005-PA4929	20050506
US 20060128722	A1	20060615	US 2005-534945	20050506
LV 13359	B	20060320	LV 2005-68	20050607
PRIORITY APPLN. INFO.:			EP 2002-79648	A 20021107
			US 2002-424579P	P 20021107
			WO 2003-EP50783	W 20031103

OTHER SOURCE(S): MARPAT 140:423584
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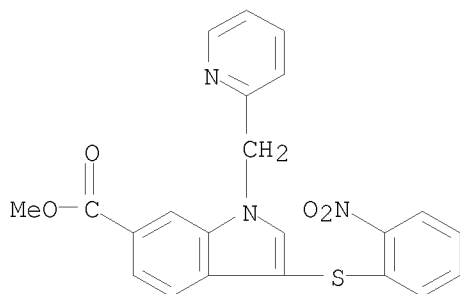


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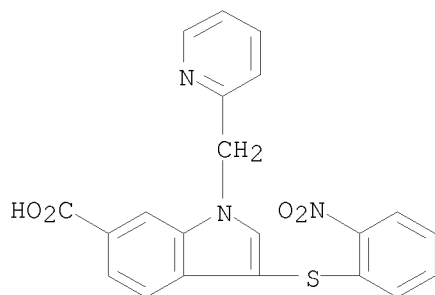


II

- AB The invention relates to a preparation of indole derivs. of formula I [wherein: X = S, S(O), SO₂; R₁ is (un)substituted 5- or 6-membered monocyclic, (hetero/homo)cyclic ring; R₂ is 2-O₂NC₆H₄, 2-cyanophenyl, 2-hydroxymethylphenyl, pyridin-2-yl, pyridin-2-yl-N-oxide, etc.; R₃ is H, halogen or C₁-4alkyl; R₄ is H, OH, C₁-4alkoxy, or halogen; R₅ is H, OH, C₁-4alkoxy, NH₂, CN, halogen, C₁-4fluoroalkyl, or NO₂, etc.], useful for the treatment of androgen-receptor related diseases. Anti-androgenic activity of the invented compds. was determined in an in vitro bioassay of Chinese hamster ovary (CHO) cells stably transfected with the human androgen receptor expression plasmid and a reporter plasmid in which the MMTV-promoter was linked to the luciferase reporter gene. For instance, indole derivs. II (EC₅₀ < 5 nM; efficacy > 0.8) was prepared via N-benylation of 6-methoxyindole by 3,5-difluorobenzyl bromide, and subsequent addition of the obtained 1-(3,5-difluorobenzyl)-6-methoxy-1H-indole to 2-nitrobenzenesulfonyl chloride (example 1).
- IT 691400-54-1P 691400-55-2P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (intermediate; preparation of indole derivs. useful in the treatment of androgen-receptor related diseases)
- RN 691400-54-1 CAPLUS
- CN 1H-Indole-6-carboxylic acid, 3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)-, methyl ester (CA INDEX NAME)



- RN 691400-55-2 CAPLUS
- CN 1H-Indole-6-carboxylic acid, 3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)- (CA INDEX NAME)



- IT 691399-74-3P 691400-66-5P
 RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP

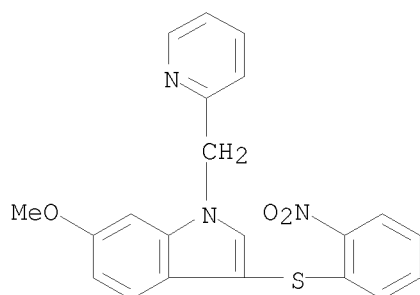
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(Preparation); RACT (Reactant or reagent); USES (Uses)

(preparation of indole derivs. useful in the treatment of androgen-receptor related diseases)

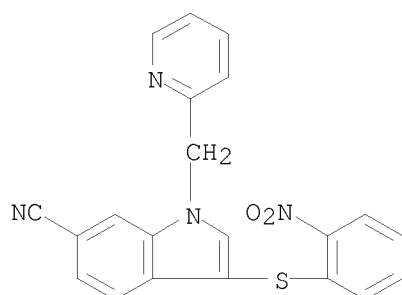
RN 691399-74-3 CAPLUS

CN 1H-Indole, 6-methoxy-3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)- (CA INDEX NAME)



RN 691400-66-5 CAPLUS

CN 1H-Indole-6-carbonitrile, 3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)- (CA INDEX NAME)



IT 691400-22-3P 691400-35-8P 691400-37-0P
691400-38-1P 691400-52-9P 691400-56-3P
691400-57-4P 691400-73-4P

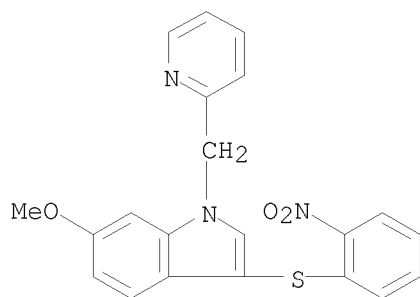
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of indole derivs. useful in the treatment of androgen-receptor related diseases)

RN 691400-22-3 CAPLUS

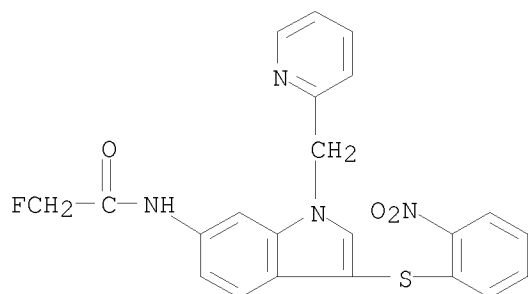
CN 1H-Indole, 6-methoxy-3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)-, hydrochloride (1:1) (CA INDEX NAME)

10/534,945

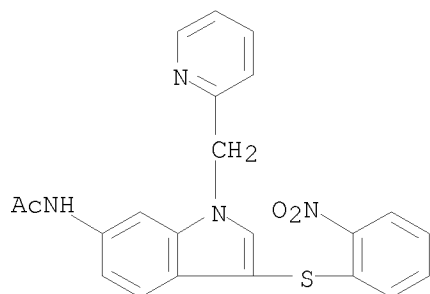


● HCl

RN 691400-35-8 CAPLUS
CN Acetamide, 2-fluoro-N-[3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)-1H-indol-6-yl]- (CA INDEX NAME)

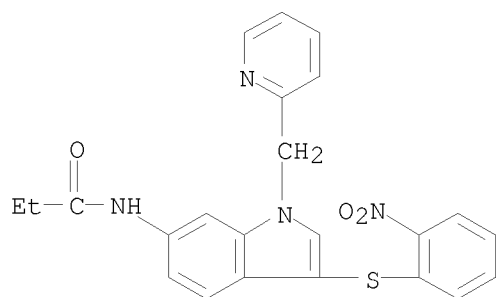


RN 691400-37-0 CAPLUS
CN Acetamide, N-[3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)-1H-indol-6-yl]- (CA INDEX NAME)

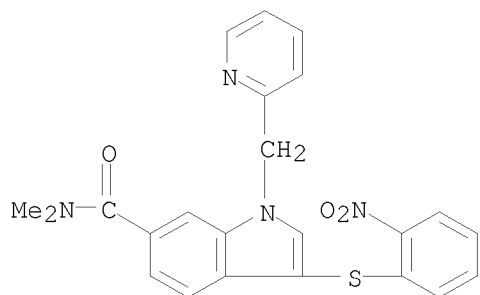


RN 691400-38-1 CAPLUS
CN Propanamide, N-[3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)-1H-indol-6-yl]- (CA INDEX NAME)

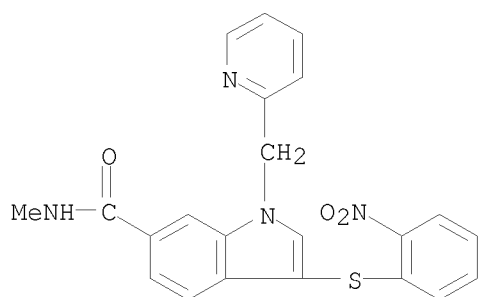
10/534,945



RN 691400-52-9 CAPLUS
CN 1H-Indole-6-carboxamide, N,N-dimethyl-3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)- (CA INDEX NAME)

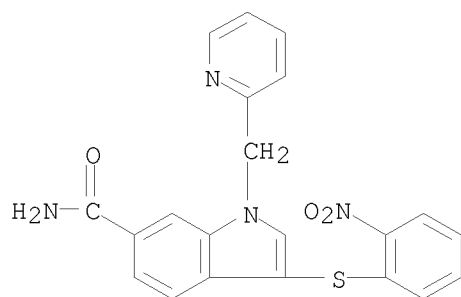


RN 691400-56-3 CAPLUS
CN 1H-Indole-6-carboxamide, N-methyl-3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)- (CA INDEX NAME)

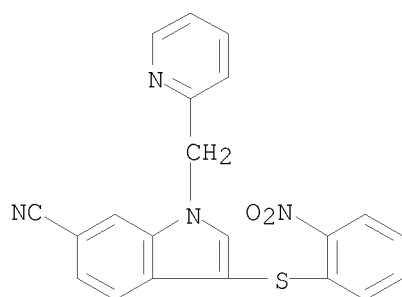


RN 691400-57-4 CAPLUS
CN 1H-Indole-6-carboxamide, 3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)- (CA INDEX NAME)

10/534,945



RN 691400-73-4 CAPLUS
CN 1H-Indole-6-carbonitrile, 3-[(2-nitrophenyl)thio]-1-(2-pyridinylmethyl)-,
hydrochloride (1:1) (CA INDEX NAME)



● HCl

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